

(12)

(21) 2 491 930

(22) 02.07.2003

(51) Int. Cl. 7: **C08G 73/18**

(85) 05.01.2005

(86) PCT/EP03/007020

(87) WO04/005373

(30) 102 30 477.7 DE 06.07.2002

(71) PEMEAS GMBH,  
65926, FRANKFURT AM MAIN, XX (DE).

(72) SINIGERSKY, VESSELIN (BG).

KLAPPER, MARKUS (DE).  
MUELLEN, KLAUS (DE).  
WU, WEICHENG (DE).  
BAURMEISTER, JOCHEN (DE).  
UENSAL, OEMER (DE).

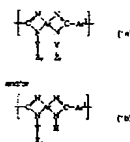
(74) BERESKIN & PARR

(54) POLYAZOLES FONCTIONNALISEES, LEUR PROCEDE DE PRODUCTION ET LEUR UTILISATION

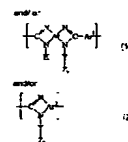
(54) FUNCTIONALIZED POLYAZOLES, METHOD FOR THE PRODUCTION THEREOF, AND USE THEREOF

(57)

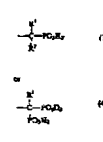
The invention relates to functionalized polyazoles containing recurrent imidazole units of general formula (1a) and/or (1b) and/or (1c) and/or (2), in which the radicals Ar, Ar1 and Ar2 represent tetravalent, divalent, or trivalent aromatic or heteroaromatic groups, Y represents a bond or a group comprising 1 to 20 carbon atoms, v represents a whole number between 1 and 10, and Z represents a group of general formula (3) or (4), in which R1 and R2 independently represent a carbon atom or a group comprising 1 to 20 carbon atoms. The inventive functionalized polyazoles are characterized by the fact that the polyazole is soluble in organic solvents. The invention also relates to intermediate products and methods for producing said polyazoles as well as polymer electrolyte membranes and fuel cells containing the inventive polyazoles.



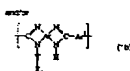
(1a)



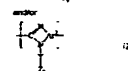
(1b)



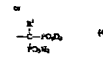
(1c)



(2)



(3)



(4)



Office de la Propriété  
Intellectuelle  
du Canada

Un organisme  
d'Industrie Canada

Canadian  
Intellectual Property  
Office

An agency of  
Industry Canada

CA 2491930 A1 2004/01/15

(21) **2 491 930**

(12) **DEMANDE DE BREVET CANADIEN  
CANADIAN PATENT APPLICATION**

(13) **A1**

(86) Date de dépôt PCT/PCT Filing Date: 2003/07/02

(87) Date publication PCT/PCT Publication Date: 2004/01/15

(85) Entrée phase nationale/National Entry: 2005/01/05

(88) N° demande PCT/PCT Application No.: EP 2003/007020

(87) N° publication PCT/PCT Publication No.: 2004/005373

(30) Priorité/Priority: 2002/07/06 (102 30 477.7) DE

(51) Cl.Int.<sup>7</sup>/Int.Cl.<sup>7</sup> C08G 73/18

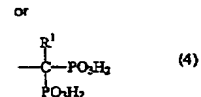
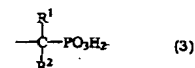
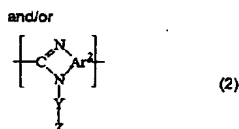
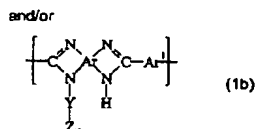
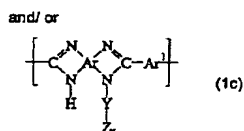
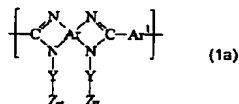
(71) Demandeur/Applicant:  
PEMEAS GMBH, DE

(72) Inventeurs/Inventors:  
MUELLEN, KLAUS, DE;  
KLAPPER, MARKUS, DE;  
SINIGERSKY, VESSELIN, BG;  
WU, WEICHENG, DE;  
UENSAL, OEMER, DE;  
BAURMEISTER, JOCHEN, DE

(74) Agent: BERESKIN & PARR

(54) Titre : POLYAZOLES FONCTIONNALISES, LEUR PROCEDE DE PRODUCTION ET LEUR UTILISATION

(54) Title: FUNCTIONALIZED POLYAZOLES, METHOD FOR THE PRODUCTION THEREOF, AND USE THEREOF



(57) Abrégé/Abstract:

The invention relates to functionalized polyazoles containing recurrent imidazole units of general formula (1a) and/or (1b) and/or (1c) and/or (2), in which the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> represent tetravalent, divalent, or trivalent aromatic or heteroaromatic groups, Y represents a bond or a group comprising 1 to 20 carbon atoms, v represents a whole number between 1 and 10, and Z represents a group of general formula (3) or (4), in which R<sup>1</sup> and R<sup>2</sup> independently represent a carbon atom or a group comprising 1 to 20 carbon atoms. The inventive functionalized polyazoles are characterized by the fact that the polyazole is soluble in organic solvents. The invention also relates to intermediate products and methods for producing said polyazoles as well as polymer electrolyte membranes and fuel cells containing the inventive polyazoles.

Canada

<http://opic.gc.ca> • Ottawa-Hull K1A 0C9 • <http://cipo.gc.ca>

OPIC • CIP0 191

OPIC



CIP0

## Abstract

Functionalized polyazoles, process for preparing them and their use

The present invention relates to functionalized polyazoles comprising recurring imidazole units of the general formula



and/or



and/or



and/or



where the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> are tetravalent, divalent or trivalent aromatic or heteroaromatic groups,

Y is a bond or a group having from 1 to 20 carbon atoms,

v is an integer from 1 to 10 and

Z is a group of the general formula



or



where R<sup>1</sup> and R<sup>2</sup> are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms, characterized in that the polyazole is soluble in organic solvents. Furthermore, the present invention relates to intermediates and to a process for preparing the polyazoles. Furthermore, the present invention describes polymer electrolyte membranes and fuel cells in which the polyazoles of the invention are present.

## Functionalized polyazoles, process for preparing them and their use

The present invention relates to polymers based on polyazoles and functionalized with phosphonic acid groups, with, owing to their excellent chemical and thermal properties, can be used for a variety of purposes and are particularly suitable for use in polymer electrolyte membrane (PEM) fuel cells.

Polyazoles such as polybenzimidazoles (@Celazole) have been known for a long time. Such polybenzimidazoles (PBIs) are usually prepared by reacting 3,3',4,4'-tetraaminobiphenyl with isophthalic acid or diphenylisophthalic acid or esters thereof in a solid-state polymerization. The resulting prepolymer solidifies in the reactor and is subsequently comminuted mechanically. The pulverulent prepolymer is subsequently fully polymerized at temperatures of up to 400°C and the desired polybenzimidazole is obtained.

To produce polymer films, the PBI is dissolved in polar, aprotic solvents such as dimethylacetamide (DMAc) and a film is produced by classical methods.

Proton-conducting polyazole membranes, i.e. polyazole membranes doped with acid, for use in PEM fuel cells are already known. Thus, J. Electrochem. Soc., Volume 142, No. 7, 1995, pp. L121-L123, describes doping a polybenzimidazole in phosphoric acid. Here, the basic polyazole film is doped with concentrated phosphoric acid or sulfuric acid and subsequently acts as proton conductor and separator in polymer electrolyte membrane fuel cells (PEM fuel cells).

Due to the excellent properties of the polyazole polymer, such polymer electrolyte membranes can, when processed to produce membrane-electrode units (MEUs), be used in fuel cells at long-term operating temperatures above 100°C, in particular above 120°C. This high long-term operating temperature allows the activity of the catalysts based on noble metals which are present in the membrane-electrode unit (MEU) to be increased. Particularly when using reformates of hydrocarbons, significant amounts of carbon monoxide are present in the reformer gas and these usually have to be removed by means of a costly gas work-up or gas purification. The ability to increase the operating temperature enables significantly higher concentrations of CO impurities to be tolerated over the long term.

The use of polymer electrolyte membranes based on polyazole polymers allows, firstly, the costly gas work-up or gas purification to be omitted in some cases and, secondly, the catalyst loading in the membrane-electrode unit to be reduced. Both

are indispensable prerequisites for large-scale use of PEM fuel cells, since otherwise the costs of a PEM fuel cell system are too high.

Prog. Polym. Sci. 25 (2000) 1463-1502 discloses a C<sub>2</sub>-alkylphosphonated polybenzimidazole, i.e. a polybenzimidazole having covalent bound phosphonic acid groups. The conductivity observed is 10<sup>-3</sup> S/cm, measured as a press compact. Although the polymer described has an elevated conductivity, it is unsuitable for the formation of polymer films by casting from polar, aprotic solvents such as dimethylacetamide (DMAc). The polymer obtained by deprotonation of polybenzimidazole with lithium hydride and subsequent reaction with 2-chloroethylphosphonic acid is, according to Rikukawa et al., insoluble in all organic solvents and therefore unsuitable for the production of polymer membranes by classical methods.

The authors assume that this is attributable to crosslinking of the polymer as a result of aggregation of the phosphonate groups during the substitution reaction.

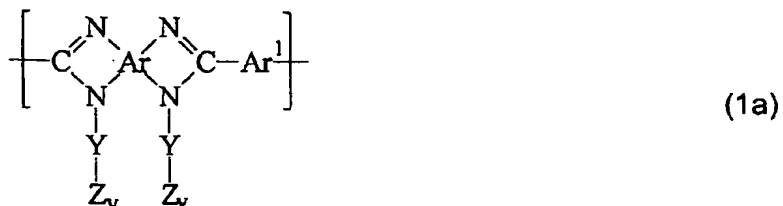
A problem with known PEM fuel cells is that their performance decreases with increasing operating time. Furthermore, the efficiency of fuel cells in respect of the conversion of hydrogen needs to be improved.

It is therefore an object of the present invention to provide improved fuel cells which have a longer life and also better performance. Furthermore, the hydrogen conversion of the fuel cell should be increased.

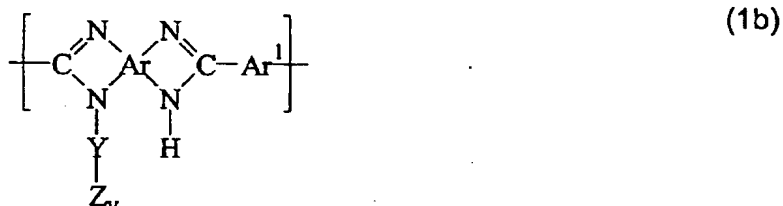
The fuel cells should be able to be used, in particular, at operating temperatures above 100°C and make do without additional humidification of the fuel gas.

Furthermore, ionomers are to be made available for use in high-temperature fuel cells, and the ionomers should be suitable, in particular, for polyazole membranes doped with phosphoric acid.

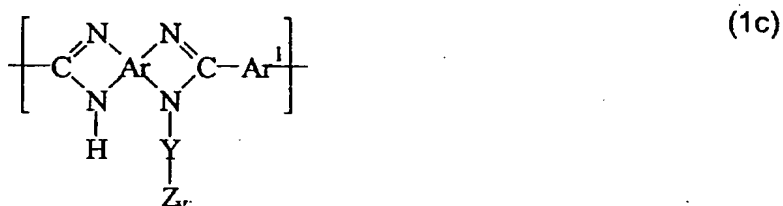
The objects mentioned above are achieved by polyazoles which are soluble in organic solvents and comprise the recurring structural units of the general formulae (1a), (1b), (1c)



and/or



and/or



and/or of the general formula (2)



where the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> are tetravalent, divalent or trivalent aromatic or heteroaromatic groups,

Y is a bond or a group having from 1 to 20 carbon atoms,

v is the number of groups Z which are bound to the group Y and is an integer from 1 to 10, and

Z is a group of the general formula (3)



or of the general formula (4)



where R1 and R2 are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms.

5 The group Y is preferably a bond or a group having 1 or 2 carbon atoms.

This is particularly surprising to a person skilled in the art because it has hitherto had to be assumed, on the base of the disclosure of Rikukawa et al., that such a polymer would be insoluble in organic solvents as a result of the aggregation of phosphonate  
10 groups. For the purposes of the invention, soluble means that at least 0.1 gram of the polymer of the invention can be dissolved in 100 gram of N,N-dimethylacetamide at 100°C.

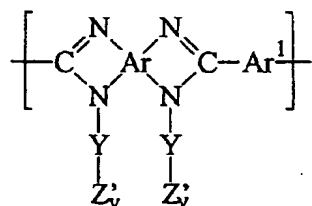
Solutions are usually homogeneous mixtures which can be employed for producing  
15 films. Insoluble material which may occur on dissolution can be separated off in a known manner.

The polymers of the invention have a number of further advantages. In particular, the polymers containing phosphonic acid groups have a high conductivity of at least  
20 0.01 S/cm, in particular at least 0.02 S/cm, at 120°C. These values are determined by impedance spectroscopy.

The polymers containing phosphonic acid groups can be obtained, inter alia, by hydrolysis of polyazoles which have been modified with phosphonic ester groups.  
25 Such polymers are valuable intermediates which are likewise provided by the invention.

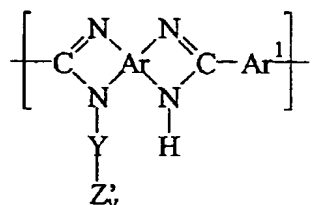
These functionalized polyazoles comprise recurring imidazole units of the general formula





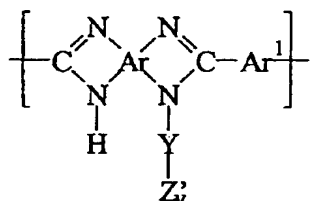
(1a')

and/or



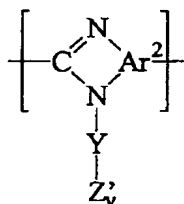
(1b')

and/or



(1c')

and/or



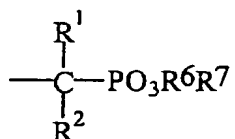
(2')

where the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> are tetravalent, divalent or trivalent aromatic or heteroaromatic groups,

Y is a bond or a group having from 1 to 20 carbon atoms,

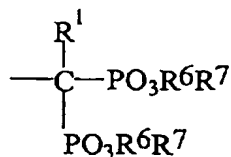
v is the number of groups Z which are bound to the group Y and is an integer from 1 to 10 and

Z' is a group of the general formula



(3')

or



(4').

where  $\text{R}^1$  and  $\text{R}^2$  are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms and  $\text{R}^6$  and  $\text{R}^7$  are each, independently of one another, a group having from 1 to 20 carbon atoms.

5 The radicals Ar, Ar1 and Ar2 of the above formulae are tetravalent, divalent or trivalent aromatic or heteroaromatic groups which may be monocyclic or polycyclic. According to the invention, aromatic groups are radicals of monocyclic or polycyclic aromatic compounds which preferably have from 6 to 14, in particular from 6 to 12, carbon atoms. Heteroaromatic groups are aryl radicals in which at least one CH  
10 group has been replaced by N and/or at least two adjacent CH groups have been replaced by S, NH or O. Aromatic or heteroaromatic groups which are preferred for the purposes of the invention are derived from benzene, naphthalene, biphenyl, diphenyl ether, diphenylmethane, diphenyldimethylmethane, bisphenone, diphenyl sulfone, thiophene, furan, pyrrole, thiazole, oxazole, imidazole, isothiazole, isoxazole,  
15 pyrazole, 1,3,4-oxadiazole, 2,5-diphenyl-1,3,4-oxadiazole, 1,3,4-thiadiazole, 1,3,4-triazole, 2,5-diphenyl-1,3,4-triazole, 1,2,5-triphenyl-1,3,4-triazole, 1,2,4-oxadiazole, 1,2,4-thiadiazole, 1,2,4-triazole, 1,2,3-triazole, 1,2,3,4-tetrazole, benzo[b]thiophene, benzo[b]furan, indole, benzo[c]thiophene, benzo[c]furan, isoindole, benzoxazole, benzothiazole, benzimidazole, benzisoxazole, benzisothiazole, benzopyrazole,  
20 benzothiadiazole, benzotriazole, dibenzofuran, dibenzothiophene, carbazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,3,5-triazine, 1,2,4-triazine, 1,2,4,5-triazine, quinoline, isoquinoline, quinoxaline, quinazoline, cinnoline, 1,8-naphthyridine, 1,5-naphthyridine, 1,6-naphthyridine, 1,7-naphthyridine, phthalazine, pyridopyrimidine, purine, pteridine or 4H-quinolizine, diphenyl ether, anthracene and  
25 phenanthrene.

The aromatic and/or heteroaromatic groups may be substituted. Preferred substituents include halogens, groups having from 1 to 20 carbon atoms, nitro, sulfonic acid, sulfonic ester, sulfinic acid, sulfinic ester, thiol, cyanide, hydroxy groups  
30 and also groups of the general formulae  $\text{NR}^8\text{R}^9$  and  $\text{N}^+\text{R}^8\text{R}^9\text{R}^{10}$ , where the radicals  $\text{R}^8$ ,  $\text{R}^9$  and  $\text{R}^{10}$  are each, independently of one another, hydrogen, an alkyl group having from 1 to 6 carbon atoms, an alkoxy group having from 1 to 6 carbon atoms

or an aryl group. According to the invention, the expression "halogen" refers to a fluorine, chlorine, bromine or iodine atom.

The expression "group having from 1 to 20 carbon atoms" refers to radicals of organic compounds having from 1 to 20 carbon atoms. Apart from the aromatic and heteroaromatic groups mentioned above, it encompasses, inter alia, alkyl, cycloalkyl, alkoxy, alkanoyl, alkoxycarbonyl groups, alkenyl groups having from 2 to 20 carbon atoms and also cycloalkoxy and cycloalkylthio groups having from 3 to 20 carbon atoms and heteroaliphatic groups which comprise, in particular, oxygen, nitrogen, sulfur and phosphorus atoms in addition to carbon and hydrogen atoms. The groups mentioned can be branched or unbranched.

Preferred alkyl groups include the methyl, ethyl, propyl, isopropyl, 1-butyl, 2-butyl, 2-methylpropyl, tert-butyl, pentyl, 2-methylbutyl, 1,1-dimethylpropyl, hexyl, heptyl, octyl, 1,1,3,3-tetramethylbutyl, nonyl, 1-decyl, 2-decyl, undecyl, dodecyl, pentadecyl and eicosyl group.

Preferred cycloalkyl groups include the cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl groups which may each be substituted by branched or unbranched alkyl groups.

Preferred alkenyl groups include the vinyl, allyl, 2-methyl-2-propenyl, 2-butenyl, 2-pentenyl, 2-decenyl and 2-eicosenyl group.

Preferred alkynyl groups include the ethynyl, propargyl, 2-methyl-2-propynyl, 2-butylnyl, 2-pentylnyl and 2-decylnyl group.

Preferred alkanoyl groups include the formyl, acetyl, propionyl, 2-methylpropionyl, butyryl, valeroyl, pivaloyl, hexanoyl, decanoyl and dodecanoyl group.

Preferred alkoxycarbonyl groups include the methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, butoxycarbonyl, tert-butoxycarbonyl, hexyloxycarbonyl, 2-methylhexyloxycarbonyl, decyloxycarbonyl or dodecyloxycarbonyl group.

Preferred alkoxy groups include alkoxy groups whose hydrocarbon radical is one of the abovementioned preferred alkyl groups.

Preferred cycloalkoxy groups include cycloalkoxy groups whose hydrocarbon radical is one of the abovementioned preferred cycloalkyl groups.

Preferred heteroaliphatic groups include the abovementioned preferred cycloalkyl radicals in which at least one carbon unit has been replaced by O, S or an  $\text{NR}^8$  group, where  $\text{R}^8$  is hydrogen, an alkyl group having from 1 to 6 carbon atoms, an alkoxy group having from 1 to 6 carbon atoms or an aryl group.

According to the invention, very particular preference is given to branched or unbranched alkyl or alkoxy groups having from 1 to 20 carbon atoms, preferably from 1 to 12 carbon atoms, advantageously from 1 to 6 carbon atoms, in particular from 1 to 4 carbon atoms, and cycloalkyl or cycloalkyloxy groups having from 3 to 20 carbon atoms, preferably from 5 to 6 carbon atoms.

Although one or more hydrogen atoms in the abovementioned radicals can be replaced by the abovementioned halogen atoms, preferably chlorine or fluorine, thiol or hydroxy groups or groups of the general formulae  $\text{NR}^8\text{R}^9$  and  $\text{N}^+\text{R}^8\text{R}^9\text{R}^{10}$ , where the radicals  $\text{R}^8$ ,  $\text{R}^9$  and  $\text{R}^{10}$  are each, independently of one another, hydrogen, an alkyl group having from 1 to 6 carbon atoms, an alkoxy group having from 1 to 6 carbon atoms or an aryl group, unsubstituted radicals have been found to be particularly advantageous.

For the purposes of the present invention,  $\text{Ar}^1$  can have any substitution pattern; in the case of phenylene, for example,  $\text{Ar}^1$  can be ortho-, meta- or para-phenylene. Particularly preferred groups  $\text{Ar}$ ,  $\text{Ar}^1$  and  $\text{Ar}^2$  are derived from benzene and biphenylene, which may also be substituted.

The polymer of the invention can further comprise recurring azole units of the general formula ("copolymer")



or/and



The radicals X are an oxygen atom (benzoxal unit), a sulfur atom (benzothiazole unit) or an amino group (benzimidazole unit) which may bear a group having 1-20 carbon atoms, preferably a branched or unbranched alkyl, cycloalkyl or alkoxy group or a substituted or unsubstituted aryl or heteroaryl group, as further radical.

If polymers having recurring azole units of the formula (12) are used for the purposes of the present invention, the radicals X within a recurring unit are, according to a preferred embodiment, identical.

Although a polymer which can be used according to the invention can in principle have different recurring units, it preferably has only identical recurring units, without this constituting a restriction.

According to a particular aspect of the present invention, the molar ratio of phosphorus to nitrogen, viz. the  $n(P)/n(N)$  ratio, measured by means of elemental analysis is from 0.02 to 0.5, preferably from 0.05 to 0.35 and very particularly preferably from 0.07 to 0.25, without this constituting a restriction.

The polyazoles modified with phosphonate groups can be prepared by a process in which polymers comprising recurring imidazole units of the general formula



or

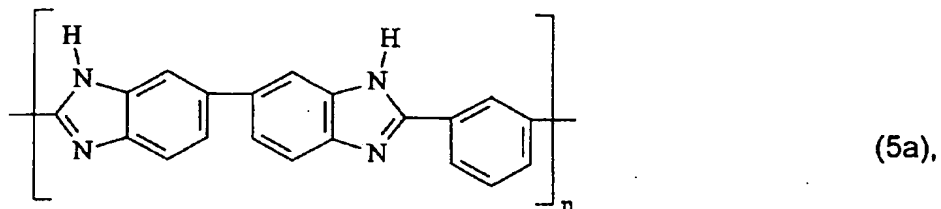


are dissolved in a solvent in a step A).

In a further preferred embodiment of the present invention, a copolymer comprising at least two units of the formula (5) and/or (6) which differ from one another is used as polymer comprising recurring imidazole units. Furthermore, these copolymers can further comprise groups of the formulae (12) and/or (13).

The number of recurring imidazole units (5) and/or (6) in the polymer used is preferably greater than or equal to 10. Particularly preferred polymers comprise at least 100 recurring imidazole units (5) and/or (6).

For the purposes of the present invention, preference is given to using polymers comprising recurring benzimidazole units. An example of an extremely advantageous polymer comprising recurring benzimidazole units has the formula (5a):



where  $n$  is an integer greater than or equal to 10, preferably greater than or equal to 100.

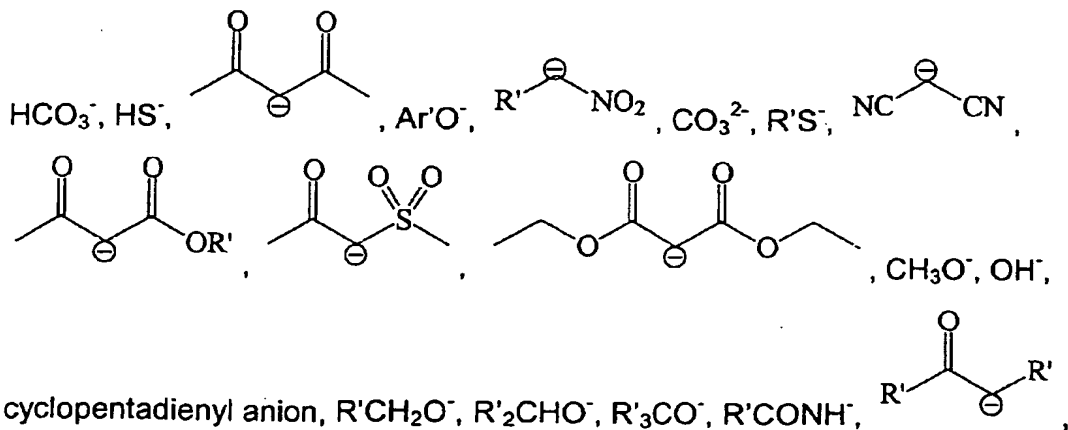
Solvents include, in particular, polar organic solvents, in particular N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide and N-methyl-2-pyrrolidone, preferably N,N-dimethylacetamide.

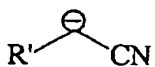
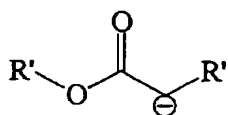
The polyazoles preferably have a high molecular weight. Measured as intrinsic viscosity, this is preferably at least 1.0 dl/g and particularly preferably at least 1.3 dl/g, measured at 25°C in DMAc, without this constituting a restriction.

To measure the intrinsic viscosity, 0.4 g of the polyazole according to the invention is dissolved in 60 ml of 98% sulfuric acid at 80°C. The intrinsic viscosity of this solution is determined at 25°C using an Ubbelohde viscometer in accordance with DIN 53728.

The reaction mixture obtained in step A) is subjected to deprotonation of the aromatic NH groups by means of a base in step B). Preferred bases have a  $pK_B$  at 25°C of less than or equal to 5, in particular less than or equal to 4, particularly preferably less than or equal to 2.

Bases which are particularly preferably used according to the invention include, inter alia,  $NH_3$ ,  $R'_3N$ ,  $R'NH_2$  and  $R'_2NH$  and also salts of the anions





,  $\text{HC}\equiv\text{C}^-$ ,  $\text{Ar}'_3\text{C}^-$ ,  $\text{Ar}'_2\text{CH}^-$ ,  $\text{H}^-$ ,  $\text{NH}_2^-$ ,  $\text{PhCH}_2^-$ , allyl anion,  $\text{Ph}^-$ ,  $\text{CH}_2=\text{CH}^-$ ,  $\text{Cyclo-C}_3\text{H}_5^-$ ,  $\text{CH}_3^-$ ,  $\text{C}_2\text{H}_5^-$ ,  $(\text{CH}_3)_2\text{CH}^-$  and  $(\text{CH}_3)_3\text{C}^-$ . In these formulae,  $\text{R}'$  is an alkyl or cycloalkyl radical having from 1 to 20 carbon atoms.  $\text{Ar}'$  is an aryl radical.  $\text{Ph}$  is a phenyl group. The salts mentioned preferably have alkali metal cations such as  $\text{Li}^+$ ,  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Rb}^+$  and  $\text{Cs}^+$ , alkaline earth metal cations such as  $\text{Be}^{2+}$ ,  $\text{Mg}^{2+}$  and  $\text{Ca}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{Zn}^{2+}$ ,  $^+\text{N}(\text{CH}_3)_4$ ,  $^+\text{N}(\text{C}_2\text{H}_5)_4$ ,  $^+\text{N}(\text{C}_3\text{H}_7)_4$ , in particular  $\text{Li}^+$  and  $\text{Na}^+$ , as counterion.  $\text{Li}_2\text{CO}_3$ ,  $\text{LiH}$  and  $\text{NaH}$ , in particular  $\text{Li}_2\text{CO}_3$ , have been found to be very particularly useful for the deprotonation.

The amount of base to be used depends on the desired degree of modification of the polymer. Preference is given to using from 0.01 to 5 molar equivalents, advantageously from 0.1 to 2.5 molar equivalents, more preferably from 0.7 to 1.5 molar equivalents, in particular from 0.9 to 1.1 molar equivalents, base, based on N-H groups in the starting polymer.

The polyazole solution prior to reaction with the base preferably has only a low water content. In a particular embodiment, the water content of the solution is 0.5% by weight or less, particularly preferably 0.3% by weight or less.

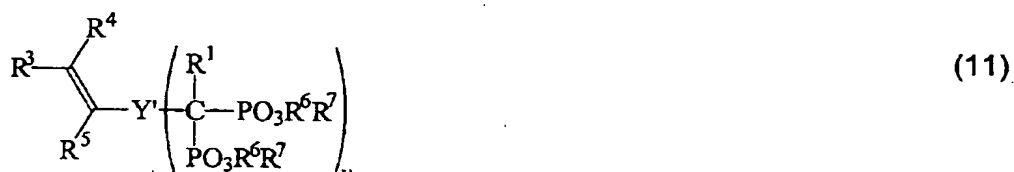
The reaction of the polymer comprising recurring imidazole units with the base can be carried out at atmospheric pressure, subatmospheric pressure or superatmospheric pressure. The reaction temperature is also not critical. However, it is generally in the range  $-20^\circ\text{C}$ - $200^\circ\text{C}$ , preferably  $20^\circ\text{C}$ - $200^\circ\text{C}$  and particularly preferably  $50^\circ\text{C}$ - $120^\circ\text{C}$ , without this constituting a restriction.

The reaction time depends on the above-described parameters. Significant evolution of gas usually occurs after only a few minutes, without this constituting a restriction. If the complete deprotonation is to be achieved, a longer reaction time which can be in the range from 10 minutes to 48 hours will sometimes be necessary. To avoid secondary reactions, the use of inert gas, in particular nitrogen and argon, has likewise been found to be particularly useful.

For the purposes of the present invention, the reaction mixture from step B) is reacted in a further step C) with at least one phosphonate of the general formulae



and/or



preferably a phosphonate of the general formula (7), (8) and/or (9). Here, the phosphonates can be used individually or as a mixture.

The radicals  $\text{R}^1$  and  $\text{R}^2$  are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms.

The radicals  $\text{R}^3$ ,  $\text{R}^4$  and  $\text{R}^5$  are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms, preferably a branched or unbranched alkyl, cycloalkyl or alkoxy group or a substituted or unsubstituted aryl or heteroaryl group. Alkyl, cycloalkyl, alkoxy, aryl or heteroaryl groups which are particularly preferred according to the invention are the abovementioned preferred radicals, as long as they have from 1 to 20, preferably from 1 to 12, advantageously from 1 to 6, in particular from 1 to 4, carbon atoms.



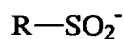
It has been found to be very particularly appropriate for at least one of the radicals  $R^3$ ,  $R^4$  and  $R^5$  to be a hydrogen atom, and preference is given to all of the radicals  $R^3$ ,  $R^4$  and  $R^5$  being hydrogen atoms.

5 The radicals  $R^6$  and  $R^7$  are each, independently of one another, a group having from 1 to 20 carbon atoms.

10 The radical Y is a bond or a group having from 1 to 20 carbon atoms, preferably a branched or unbranched alkyl, cycloalkyl or alkoxy group or a substituted or unsubstituted aryl or heteroaryl group. Alkyl, cycloalkyl, alkoxy, aryl or heteroaryl groups which are particularly preferred according to the invention include the abovementioned preferred radicals, as long as they have from 1 to 20, preferably from 1 to 12, advantageously from 1 to 6, in particular from 1 to 4, carbon atoms. The group Y is particularly preferably a group having 2 carbon atoms.

15 The radical X is a leaving group. For the purposes of the invention, leaving groups are the groups which are split off in the course of a substitution reaction, in the present case a nucleophilic substitution reaction. For further details, reference may be made to the specialist literature, for example March, Jerry, Advanced Organic Chemistry, Wiley Interscience, 1985, in particular p. 179.

20 Preferred leaving groups include the following radicals:  
halogen atoms, preferably a chlorine atom, bromine atom or iodine atom,  
groups of the general formula



25 where R is a fluorinated or unfluorinated, linear or branched aliphatic or cycloaliphatic radical or a substituted or unsubstituted aromatic radical, preferably an alkyl radical having from 1 to 6 carbon atoms,  $CF_3$ , phenyl or para-tosyl, alkylsulfates, preferably methylsulfate, ethylsulfate,  
30 carboxylates, preferably formate, acetate.

Chlorine atoms, bromine atoms and para-toluenesulfonates have been found to be particularly useful.

35 The index v indicates the number of groups Z which are bound to the group Y and is an integer in the range from 1 to 10, preferably 1.

The radical Y' is a bond or a group having from 1 to 20 carbon atoms, preferably a branched or unbranched alkyl, cycloalkyl or alkoxy group or a substituted or unsubstituted aryl or heteroaryl group. Alkyl, cycloalkyl, alkoxy, aryl or heteroaryl groups which are particularly preferred according to the invention include the  
 5 abovementioned preferred radicals, as long as they have from 1 to 20, preferably from 1 to 12, advantageously from 1 to 6, in particular from 1 to 4, carbon atoms.

Phosphonates which are very particularly advantageous for the purposes of the invention have the formulae (7a) and (8a):



10 where m is an integer from 0 to 11, i.e. 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 and 11, preferably from 1 to 6, advantageously from 2 to 4, in particular 2.

The amount of phosphonate depends on the desired degree of modification.  
 15 Preference is given to using from 0.01 to 5 molar equivalents, advantageously from 0.1 to 2.5 molar equivalents, more preferably from 0.7 to 1.5 molar equivalents, in particular from 0.9 to 1.1 molar equivalents, of phosphonate, based on N-H groups in the starting polymer.

20 The phosphonates which can be used for preparing the polymers are known to those skilled in the art.

The reaction of the deprotonated polymer comprising recurring imidazole units with the phosphonate can be carried out at atmospheric pressure, subatmospheric  
 25 pressure or superatmospheric pressure. The reaction temperature is also not critical. however, it is generally in the range from -20° to 200°C, preferably from 20° to 200°C and particularly preferably 50°-120°C, without this constituting a restriction. The reaction time depends on the above-described parameters. In general, the reaction will be concluded after a time in the range from 2 to 48 hours.

30 To avoid secondary reactions, the use of inert gas, in particular nitrogen and argon, has likewise been found to be particularly useful.

To prepare polyazoles modified with phosphonic acid groups, the reaction mixture  
 35 from step C) can be acidified with an acid. Suitable acids preferably have a pK<sub>A</sub> at

25°C of less than 6, in particular less than 5, particularly preferably less than 4. Here, the  $pK_A$  is the negative logarithm to the base 10 of the acid constant in water.

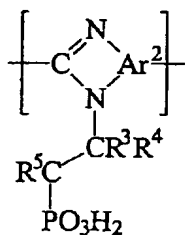
Acids which are particularly preferably used according to the invention include, inter alia,

HF-SBF<sub>5</sub>, HClO<sub>4</sub>, HI, H<sub>2</sub>SO<sub>4</sub>, HBr, HCl, Ar'SO<sub>3</sub>H, where Ar' is an aryl radical, CH(CN)<sub>3</sub>, HNO<sub>3</sub>, HF, HNO<sub>2</sub>, R'COOH, where R' is an alkyl or cycloalkyl radical having from 1 to 20 carbon atoms, HCOCH<sub>2</sub>COH, H<sub>2</sub>CO<sub>3</sub>, H<sub>2</sub>SO<sub>4</sub> and H<sub>3</sub>PO<sub>4</sub>.

The acidification of the reaction mixture from step B) can be carried out at atmospheric pressure, subatmospheric pressure, or superatmospheric pressure. The reaction temperature is also not critical. However, it is generally in the range -20°-200°C, preferably 20°-200°C and particularly preferably 25°-50°C, without this constituting a restriction.

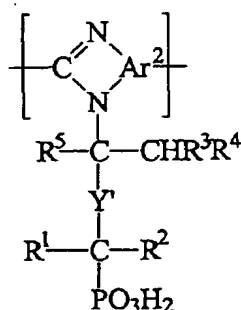
The reaction time depends on the above-described parameters. Complete reaction is generally achieved after a reaction time in the range from 10 minutes to 48 hours.

The structure of the modified polyazoles is obvious to a person skilled in the art. For example, when a phosphonate of the general formula (9) is used, the addition of the deprotonated polymer generally occurs at the carbon atom of the double bond which bears the radicals R<sup>3</sup> and R<sup>4</sup>. The reaction of a polymer comprising recurring imidazole units of the formula (6) therefore usually leads to a polymer comprising recurring imidazole units of the general formula



(2a).

In contrast, the use of a phosphonate of the general formula (8) generally results in addition onto the double bond in the opposite direction. The reaction of a polymer comprising recurring imidazole units of the formula (6) therefore usually leads to a polymer comprising recurring imidazole units of the general formula



(2b).

The isolation of the polymer containing phosphonic acid groups and comprising recurring imidazole units of the general formula (1) and/or (2) can be carried out in a manner known per se. In a preferred embodiment of the present invention, it is isolated by evaporation of the solvent at atmospheric, subatmospheric or superatmospheric pressure. In a further preferred embodiment of the present invention, it is isolated by precipitation in a nonsolvent, preferably water, alcohol or a mixture thereof, which is used in an appropriate excess.

Furthermore, the polymer of the invention has good solubility in polar solvents. This is a great advantage in further processing, because it can also be processed further from solution. The solubility of the polymer containing phosphonic acid groups in N,N-dimethylacetamide at 100°C is preferably at least 0.1 g, advantageously at least 1 g, in particular at least 5 g, in each case based on 100 g of solution.

To increase the conductivity further, the polymer of the invention containing phosphonic acid groups can advantageously be doped. Doping, i.e. the presence of dopants in the polymer, increases the proton conductivity compared to the undoped polymer. Dopants for the polymers of the invention are acids, preferably inorganic acids. For the present purposes, acids are all known Lewis and Brønsted acids, preferably inorganic Lewis and Brønsted acids. The use of polyacids, in particular isopolyacids and heteropolyacids, and also mixtures of various acids is also possible.

The conductivity of the polymer of the invention can be influenced via the degree of doping. The conductivity increases with rising concentration of dopant until a maximum value has been reached. According to the invention, the degree of doping is reported as mole of acid per mole of repeating unit of the polymer. For the purposes of the present invention, a degree of doping of from 3 to 30, in particular from 5 to 18, is preferred.

Dopants which are particularly preferred according to the invention are sulfuric acid and phosphoric acid. A very particularly preferred dopant is phosphoric acid ( $\text{H}_3\text{PO}_4$ ). This acid can be used in a concentration of less than or equal to 85% by weight, preferably in a range from 30 to 83% by weight, particularly preferably in a range from 50 to 80% by weight and very particularly preferably in a range from 60 to 75% by weight.

Conventional polyazole films are usually doped with a phosphoric acid having a concentration of at least 85% by weight. However, a problem here is that a phosphoric acid having such a high concentration can lead to a decrease in the mechanical stability of the membrane, so that this membrane fails prematurely in a fuel cell.

However, it has surprisingly been found that polyazole modified with phosphonic acid displays a significantly better swelling behavior than conventional polyazole, with the swelling increasing with increasing molar P/N ratio.

The polyazole modified with phosphonic acid groups is particularly preferably used as ionomer. The ionomer is used, in particular, for the treatment of the electrode which has been doped with a catalyst. Suitable catalysts comprise, inter alia, noble metals such as platinum and/or ruthenium. The use of an ionomer according to the invention improves the contact between the catalyst and the PEM and the performance and the long-term stability of the fuel cell is surprisingly increased.

Furthermore, a polyazole modified with phosphonic acid groups is suitable, in particular, for producing membranes for high-temperature fuel cells. For this purpose, use is made, in particular, of polyazoles whose molar P/N ratio is in the range from 0.02 to 0.25, preferably from 0.05 to 0.21, particularly preferably from 0.07 to 0.17. It has surprisingly been found that such a polyazole has a very high conductivity combined with a relatively high stability after doping with phosphoric acid having a concentration in the abovementioned ranges, so that doped polyazole films are especially suitable for use as membranes in fuel cells.

Owing to its property profile, the polymer of the invention is also suitable for application as coating to a known polyelectrolyte membrane or can be employed as a blend with known polymers, in particular polyazoles, for producing polyelectrolyte membranes. A conventional polyelectrolyte membrane can be provided on either one side or both sides with a layer of modified polyazoles, with the thickness of a polyazole coating generally being in the range from 5 to 30  $\mu\text{m}$ , preferably from 10 to

25  $\mu\text{m}$ . The thickness of the polyelectrolyte membrane is generally in the range from 5  $\mu\text{m}$  to 2000  $\mu\text{m}$ , preferably from 10  $\mu\text{m}$  to 1000  $\mu\text{m}$ , particularly preferably from 30  $\mu\text{m}$  to 200  $\mu\text{m}$ , without this constituting a restriction.

5 The present invention also provides a membrane-electrode unit comprising at least one polymer containing phosphonic acid groups according to the invention. For further information on membrane-electrode units, reference may be made to the specialist literature, in particular the patents US 4,191,618, US 4,212,714 and US 4,333,805, whose disclosure is explicitly incorporated by reference into the present  
10 patent application.

Possible fields of use for the polymers of the invention which have been modified with phosphonic acid groups include, inter alia, the use in fuel cells, in electrolysis, in capacitors and in battery systems. The polymer of the invention is particularly  
15 preferably used in fuel cells. Accordingly, fuel cells comprising the polyazoles which have been modified according to the invention are also provided by the present invention.

20 The invention is illustrated below by means of examples and comparative examples, without the invention being restricted to these examples.

The conductivity of the membrane depends greatly on the content of acid groups expressed by the ion exchange capacity (IEC). To measure the ion exchange capacity, a specimen having a diameter of 3 cm is stamped out and placed in a glass  
25 beaker containing 100 ml of water. The acid liberated is titrated with 0.1 M NaOH. The specimen is subsequently taken out, excess water is dabbed off and the specimen is dried at 160°C for 4 hours. The dry weight,  $m_0$ , is then determined gravimetrically to a precision of 0.1 mg. The ion exchange capacity is then calculated from the consumption of 0.1M NaOH to the first titration end point,  $V_1$  in ml, and the  
30 dry weight,  $m_0$  in mg, according to the following formula:

$$\text{IEC} = V_1 \cdot 300 / m_0$$

35 The swelling of membranes is calculated as the increase in area after doping with acid.

The specific conductivity is measured by means of impedance spectroscopy in a 4-pole arrangement in the potentiostatic mode using platinum electrodes (wire, 0.25 mm diameter). The distance between the current-collecting electrodes is 2 cm. The

spectrum obtained is evaluated by means of a simple model consisting of a parallel arrangement of an ohmic resistance and a capacitor. The specimen cross section of the membrane doped with phosphoric acid is measured immediately before installing the specimen. To measure the temperature dependence, the measurement cell is brought to the desired temperature in an oven and the temperature is regulated by means of a Pt-100 resistance thermometer positioned in the immediate vicinity of the specimen. After the desired temperature has been reached, the specimen is maintained at this temperature for 10 minutes before commencement of the measurement.

General procedure for the phosphoration of PBI:

80 g of a 15% strength polybenzimidazole solution in N,N-dimethylacetamide (DMAc) were slowly heated to 80°C and diluted to 350 ml with DMAc.

2.1 equivalents per element unit of the appropriate base, e.g. 11.2 g of  $K_2CO_3$  for Example 3, were then added, as a result of which the solution became deep red in color. The mixture was stirred at 80°C for 3 hours. Two equivalents per element unit of 2-bromoethylphosphonate (19.0 g) were then added dropwise to the reaction mixture from a dropping funnel. The reaction mixture was stirred at 85°C for 60 hours. After acidification, the reaction mixture was poured into three liters of a water/ethanol mixture (1:1). The precipitate was filtered off, washed thoroughly with water and methanol and dried. The polymer was dissolved in DMAc, reprecipitated in methanol and dried under reduced pressure for at least 24 hours.

Specimen	Base	Reaction time [h]	n(P)/n(N)
1	$K_2CO_3$	36	0.065
2	$K_2CO_3$	50	0.1
3	$K_2CO_3$	60	0.155
4	NaH	48	0.205
5	NaH	60	0.255
6	NaH	72	0.28
7	$K_2CO_3$	72	0.245

Properties of specimen 1-5 after doping in 85% strength  $H_3PO_4$ :

Specimen	n(P)/n(N)	Swelling in 85% $H_3PO_4$	IEC [meq/g]
1	0.065	24%	112.41
2	0.1	110%	205.14

3	0.155	not measurable	245.7
4	0.205	not measurable	356.6
5	0.255	not measurable	specimen is dissolved

Conductivity results for specimen 3 after doping with different concentrations of  $\text{H}_3\text{PO}_4$ :

	50% $\text{H}_3\text{PO}_4$	60% $\text{H}_3\text{PO}_4$	70% $\text{H}_3\text{PO}_4$	80% $\text{H}_3\text{PO}_4$
Temp. [°C]	Conductivity [S/cm]	Conductivity [S/cm]	Conductivity [S/cm]	Conductivity [S/cm]
25	0.0016	0.0033	0.016	0.084
40	0.0005	0.0008	0.008	0.056
60	0.0004	0.0009	0.007	0.045
80	0.0008	0.0017	0.010	0.051
100	0.0016	0.0037	0.018	0.064
120	0.0034	0.0075	0.029	0.077
140	0.0062	0.0122	0.034	0.080

5 Comparative examples as described by Rikukawa et al.:

A.) Model reactions

A.1) Model reaction 1

20 mg (2.4 mmol) of LiH were added to 25 ml of a solution of 0.44 g (2.3 mmol) of 2-phenylbenzimidazole in N,N-dimethylacetamide (DMAc). The mixture was heated to 85°C and stirred at this temperature for 2 hours. 0.36 g of 2-chloroethylphosphonate (90%, Aldrich) and 0.32 ml of triethylamine (dissolved in 25 ml of DMAc) were then added dropwise at 25°C over a period of 30 minutes. The reaction mixture was stirred at room temperature for 24 hours and then poured into 200 ml of water. The precipitate was filtered off and dried. The yield was 80%. A main peak was found at 366.7 g/mol in the FD mass spectrum, while the expected product has a molecular weight of 302 g/mol. An examination of the water filtrate was likewise unsuccessful.

A.2) Model reaction 2

40 mg (4.8 mmol) of LiH were added to 25 ml of a solution of 0.55 g (2.8 mmol) of 2-phenylbenzimidazole in N,N-dimethylacetamide (DMAc). The mixture was heated to 85°C and stirred at this temperature for 2 hours. 0.54 g of 2-chloroethylphosphonate (90%, Aldrich) and 1 ml of triethylamine (dissolved in 25 ml of DMAc) were then added dropwise at 25°C over a period of 30 minutes. The reaction mixture was stirred at room temperature for 24 hours and then poured into 200 ml of water. The precipitate was filtered off and dried. The yield was 85%. No main peak at 302 g/mol



was found in the FD mass spectrum. The  $^1\text{H}$ -NMR spectrum indicated that the solid was the starting component 2-phenylbenzimidazole.

#### A.3) Model reaction 3

130 mg (3.25 mmol) of NaH (60% dispersion in oil) were added to 25 ml of a solution of 0.51 g (2.6 mmol) of 2-phenylbenzimidazole in N,N-dimethylacetamide (DMAc). The mixture was heated to 85°C and stirred at this temperature for 2 hours. 0.44 g of 2-chloroethylphosphonate (90%, Aldrich) and 1 ml of triethylamine (dissolved in 25 ml of DMAc) were then added dropwise at 25°C over a period of 30 minutes. The reaction mixture was stirred at room temperature for 24 hours and then poured into 200 ml of water. The precipitate was filtered off and dried. The yield was 74%. No main peak at 302 g/mol was found in the FD mass spectrum. An examination of the water filtrate was likewise unsuccessful.

#### A.4) Model reaction 4

3 ml of triethylamine were added to 25 ml of a solution of 0.41 g (2.1 mmol) of 2-phenylbenzimidazole in N,N-dimethylacetamide (DMAc) at 25°C over a period of 30 minutes. The mixture was heated to 50°C. 0.36 g (2.2 mmol) of 2-chloroethylphosphonate (90%, Aldrich) was then added dropwise from a dropping funnel. The reaction mixture was stirred at 50°C for 24 hours and then poured into 200 ml of water. The precipitate was filtered off and dried. The yield was 85%. No main peak at 302 g/mol was found in the FD mass spectrum. An examination of the water filtrate was likewise unsuccessful.

#### B.) Polymer-analogous reaction

20 g of a 15% strength polybenzimidazole solution in N,N-dimethylacetamide (DMAc) were slowly heated to 85°C and diluted with 25 ml of DMAc. 80 mg of LiH were then added, as a result of which the solution became deep red in color. The mixture was stirred at 85°C for 3 hours. 3.12 g of 2-chloroethylphosphonate (dissolved in 25 ml of DMAc, 2.0 equivalents per repeating unit) and 3.0 ml of triethylamine were then added dropwise to the reaction mixture from a dropping funnel. The reaction mixture was stirred at 25°C for 48 hours. After acidification, the reaction mixture was poured into 1 l of a water/ethanol mixture (1:1). The dark brown precipitate was filtered off, washed thoroughly with acetone and ethanol and dried at 60°C under reduced pressure for 48 hours.

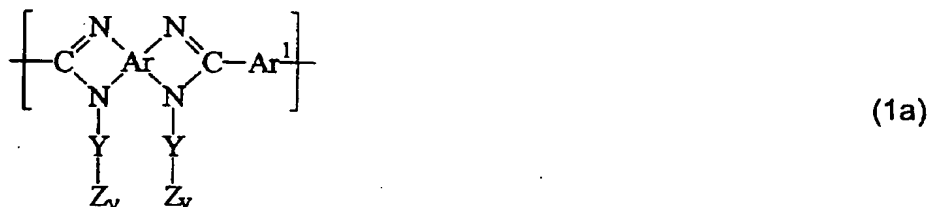
Yield: 3.7 g (53% of theory)

Solubility: Insoluble in DMSO, only soluble in concentrated sulfuric acid

Degree of modification  $n(\text{P})/n(\text{N})$ : 0.25 (4 g were obtained)

## Claims:

1. A functionalized polyazole comprising recurring imidazole units of the general formula



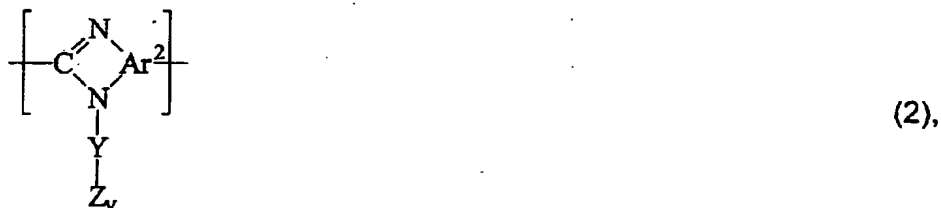
and/or



and/ or



and/or



where the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> are tetravalent, divalent or trivalent aromatic or heteroaromatic groups,

Y is a bond or a group having from 1 to 20 carbon atoms,

v is an integer from 1 to 10 and

Z is a group of the general formula



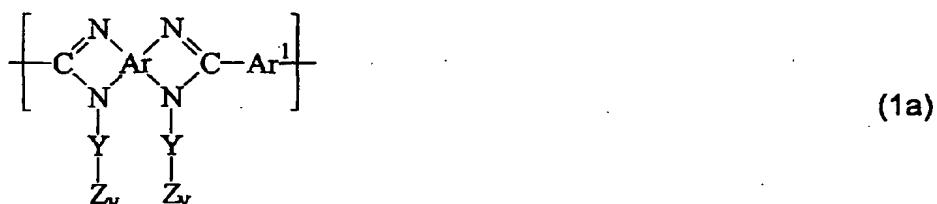
or



where  $\text{R}^1$  and  $\text{R}^2$  are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms, characterized in that the solubility of the polyazole in N,N-dimethylacetamide is at least 0.1 g, based on 100 g of solution, at 100°C.

5

2. A functionalized polyazole comprising recurring imidazole units of the general formula



and/or



and/or



and/or



where the radicals Ar, Ar<sup>1</sup> and Ar<sup>2</sup> are tetravalent, divalent or trivalent aromatic or heteroaromatic groups,

Y is a bond or a group having from 1 to 20 carbon atoms,

v is an integer from 1 to 10 and

Z' is a group of the general formula

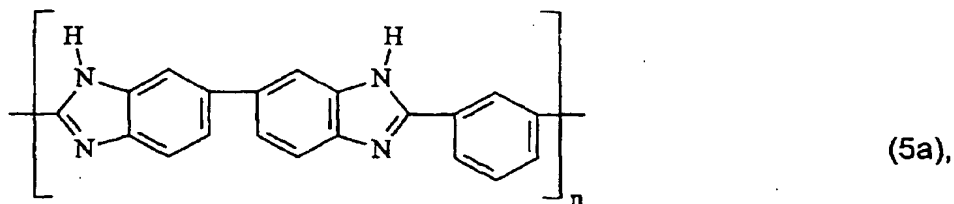


or



where R<sup>1</sup> and R<sup>2</sup> are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms and R<sup>6</sup> and R<sup>7</sup> are each, independently of one another, a group having from 1 to 20 carbon atoms.

3. The polyazole as claimed in claim 1 or 2, characterized in that the polymer comprises recurring benzimidazole units of the formula (5a):



where n is an integer greater than or equal to 10.

4. The polyazole as claimed in one or more of the preceding claims, characterized in that it is doped with an acid.
5. The polyazole as claimed in claim 4, characterized in that the degree of doping, expressed as mole of acid per mole of repeating units of the polymer, is from 3 to 15.
6. The polyazole as claimed in one or more of the preceding claims, characterized in that the group Y is a radical having 1 or 2 carbon atoms.

7. The polyazole as claimed in one or more of the preceding claims, characterized in that it has a molar ratio of phosphorus to nitrogen, P/N, measured by means of elemental analysis in the range from 0.02 to 0.5.

- 5 8. A process for preparing functionalized polyazoles as claimed in one or more of claims 2 to 7, characterized in that

A) a polymer comprising recurring imidazole units of the general formula



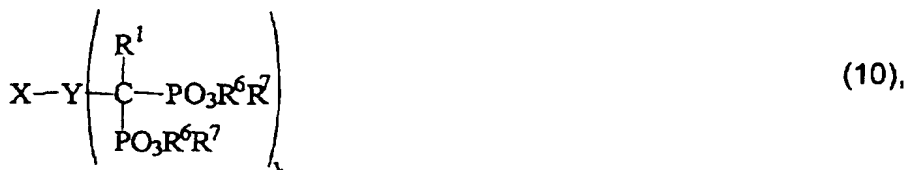
and/or



is dissolved in a solvent,

B) this solution is reacted with a base and deprotonated in this way,

10 C) the solution from step B) is reacted with at least one phosphonate of the general formulae



and/or



where  $\text{R}^3$ ,  $\text{R}^4$  and  $\text{R}^5$  are each, independently of one another, a hydrogen atom or a group having from 1 to 20 carbon atoms,

$\text{R}^6$  and  $\text{R}^7$  are each, independently of one another, a group having from 1 to 20 carbon atoms,

X is a leaving group and

Y' is a bond or a group having from 1 to 20 carbon atoms.

9. The process as claimed in claim 8 for preparing functionalized polyazoles as claimed in one or more of claims 1 and 3 to 7, characterized in that the solution resulting from C) is acidified with an acid.
10. The process as claimed in any of the preceding claims, characterized in that a base having a  $\text{pK}_a$  at 25°C of less than 7, preferably less than 6, in particular less than 5, is used in step A).
11. The process as claimed in any of the preceding claims, characterized in that phosphonates of the general formulae



where  $m$  is an integer from 0 to 11 and the radicals X,  $\text{R}^6$  and  $\text{R}^7$  are as defined above, is used as phosphonate in step B).

12. A polyazole obtainable by a process as claimed in claim 9.
13. A polymer electrolyte membrane coated with polyazoles as claimed in at least one of claims 1 to 7 and 12.
14. A polymer electrolyte membrane comprising polyazoles as claimed in at least one of claims 1 to 7 and 12.
15. A membrane-electrode unit comprising a polymer electrolyte membrane as claimed in claim 13 or 14.

16. A membrane-electrode unit comprising ionomers based on polyazoles as claimed in at least one of claims 1 to 7 and 12.
17. A fuel cell comprising a membrane-electrode unit as claimed in claim 15 or 16.